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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO	
09/412,297	10/05/1999	KANG TING	3100.006US0	9486	
33401 7	7590 11/17/2004	EXAMINE		INER	
MCDERMOTT, WILL & EMERY (LOS ANGELES OFFICE)			FORD, VANESSA L		
2049 CENTURY PARK EAST 34TH FLOOR		ART UNIT	PAPER NUMBER		
	ES, CA 90067-3208	1645			
			DATE MAILED: 11/17/2004	DATE MAILED: 11/17/2004	

Please find below and/or attached an Office communication concerning this application or proceeding.

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:	Application No.	Applicant(s)			
	09/412,297	TING, KANG			
Office Action Summary	Examiner	Art Unit			
	Vanessa L. Ford	1645			
The MAILING DATE of this communication appeared for Reply	ppears on the cover sheet with the c	correspondence address			
A SHORTENED STATUTORY PERIOD FOR REP THE MAILING DATE OF THIS COMMUNICATION - Extensions of time may be available under the provisions of 37 CFR 1 after SIX (6) MONTHS from the mailing date of this communication. - If the period for reply specified above is less than thirty (30) days, a re - If NO period for reply is specified above, the maximum statutory perio - Failure to reply within the set or extended period for reply will, by statu. Any reply received by the Office later than three months after the mail earned patent term adjustment. See 37 CFR 1.704(b).	l136(a). In no event, however, may a reply be timely within the statutory minimum of thirty (30) days d will apply and will expire SIX (6) MONTHS from the cause the application to become ABANDONE!	nely filed s will be considered timely. the mailing date of this communication. D (35 U.S.C. § 133).			
Status					
1) Responsive to communication(s) filed on 13	September 2004.				
	<u> </u>				
	Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under <i>Ex parte Quayle</i> , 1935 C.D. 11, 453 O.G. 213.				
Disposition of Claims					
4) ⊠ Claim(s) 1-53 is/are pending in the application 4a) Of the above claim(s) 3-7 and 13-50 is/are 5) □ Claim(s) is/are allowed. 6) ⊠ Claim(s) 1,2,8-12 and 51-53 is/are rejected. 7) □ Claim(s) is/are objected to. 8) □ Claim(s) are subject to restriction and/	e withdrawn from consideration.				
Application Papers					
9) ☐ The specification is objected to by the Examin 10) ☑ The drawing(s) filed on <u>05 October 1999</u> is/ard Applicant may not request that any objection to the Replacement drawing sheet(s) including the correct 11) ☐ The oath or declaration is objected to by the E	e: a)⊠ accepted or b)⊡ objected e drawing(s) be held in abeyance. See ction is required if the drawing(s) is obj	37 CFR 1.85(a). ected to. See 37 CFR 1.121(d).			
Priority under 35 U.S.C. § 119	·				
 12) Acknowledgment is made of a claim for foreign a) All b) Some * c) None of: 1. Certified copies of the priority document 2. Certified copies of the priority document 3. Copies of the certified copies of the priority application from the International Bureat * See the attached detailed Office action for a list 	ts have been received. ts have been received in Applicationity documents have been received out (PCT Rule 17.2(a)).	on No d in this National Stage			
Attachment(s)					
1) X Notice of References Cited (PTO-892)	4) 🔲 Interview Summary (
 Notice of Draftsperson's Patent Drawing Review (PTO-948) Information Disclosure Statement(s) (PTO-1449 or PTO/SB/08) 	Paper No(s)/Mail Dat 5) Notice of Informal Pa				
Paper No(s)/Mail Date <u>9/13/04</u> .	6) Other:	Company (1-10-102)			

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DETAILED ACTION

1. A request for continued examination under 37 CFR 1.114, including the fee set forth in 37 CFR 1.17(e), was filed in this application after final rejection. Since this application is eligible for continued examination under 37 CFR 1.114, and the fee set forth in 37 CFR 1.17(e) has been timely paid, the finality of the previous Office action has been withdrawn pursuant to 37 CFR 1.114. Applicant's submission filed on September 13, 2004 has been entered. Claims 1-3 have been amended. Claims 3-7 and 13-50 have been withdrawn. Claims 51-53 have been added. Applicant's amendment is acknowledged. Claims 1,2, 8-12 and 51-53 have been examined in this Office action.

Rejection Withdrawn

2. In view of Applicant's response the rejection under 35 U.S.C. 112, first paragraph of claims 1-2, 8-12 and 50 paragraph 3, pages 2-5 of the Final Office action is withdrawn.

Claim Rejections - 35 USC § 112

The following is a quotation of the first paragraph of 35 U.S.C. 112:

The specification shall contain a written description of the invention, and of the manner and process of making and using it, in such full, clear, concise, and exact terms as to enable any person skilled in the art to which it pertains, or with which it is most nearly connected, to make and use the same and shall set forth the best mode contemplated by the inventor of carrying out his invention.

3. Claims 1-2, 8-12 and 51-53 are rejected under 35 U.S.C. 112, first paragraph, because the specification, while being enabling for a method of screening for an agent

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that modulates bone mineralization, said method comprising contacting an osteogenic cell selected from the group consisting of an osteroblast, a mesenchymal, a fibroblast cell, a dura cell, a chondrocyte, a chondreoblast and a MC3T3 cell expressing a Nell-1 gene with a test agent and detecting an expression level of said Nell-1 gene in the contacted cell where a difference in the expression level of Nell-1 in the contacted cell compared to an expression level of Nell-1 in a cell that is not contacted indicates that said test agent is an agent that modulates bone mineralization does not reasonably provide enablement for claimed method of screening for an agent that modulates bone mineralization, wherein the method comprising contacting an osteogenic cell selected from the group consisting of a stem cell and a bone marrow cell. The specification does not enable any person skilled in the art to which it pertains, or with which it is most nearly connected, to make and use the invention commensurate in scope with these claims.

The specification teaches that Nell-1 expression is increased in fetal calvarial cells associated with bone formation in calvarial osteoblast-like cells in the fetus (page 42). The specification teaches that premature cranial suture closure as seen in craniosynostosis (CS) may be due to overproduction of cranial bone and therefore possibly but not definitively associated with the over-expression of the Nell-1 molecule (page 42). The specification teaches that Nell-1 mRNA was faintly expressed from day 14 of gestation with mild increase over gestation period and at day 14 is the time point when fetal calvaria starts to mineralize (page 42). The specification states "as a possible role of Nell-1, these proteins may act as a modulator interacting with other

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growth factors. The specification teaches that since $TGF\beta$ -1 is known as a regulator of osteogenesis, Nell-1's effect in enhancing mineralization may be related to its interaction with the $TGF\beta$ superfamily (page 43). The specification speculates on the function of Nell-1 based on the observation that it's expression is increased in fetal calvarial osteoblast-like cells. Although the specification teaches the use of "bone progenitor cells" which refer to any or all cells that have the capacity to ultimately form or contribute to the formation of new bone tissue (page 33).

The teachings of the cited art regarding bone progenitor cells as they relate to the claimed invention are cited below:

Bellows et al (Developmental Biology, 133, 8-13, 1989) teach that fetal or calvaria has become a standard model for bone cell metabolism (page 8). Bellows et al teach that isolated populations express osteoblastic properties comprise a heterogenous mixture of cell which include fibroblasts, chondrocytes, undifferentiated mesenchymal cells and cell at various stages of osteoblast differentiation (page 8). Bellows et al teach that precursors of osteogenic cell are believed to be derived from stem cells (page 12). Peterson et al (U. S. Patent No. 6,200,606 B1, published March 13, 2001) teach that the process of biological differentiation, which underlies the diversity of cell types exhibited by bone marrow, is the general process by which specialized, committed cell types arise from less specialized, primitive cell types (column 2). Peterson et al teach that differentiation may conveniently be thought of as a series of steps along a pathway, in which each step is occupied by a particular cell type potentially having unique genetic and phenotypic characteristics (column 2). Peterson

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et al teach that in the typical course of differentiation a pluripotent stem cell proceeds through one or more intermediate stage cellular divisions, ending ultimately in the appearance of one or more specialized cell types, such as T lymphocytes and osteocytes (column 2). Peterson et al teach that the uncommitted cell types which precede the fully differentiated forms, and which may or may not be true stem cells, are defined as precursor cells (column 2). Peterson et al teach that the precise signals that trigger differentiation down a particular path are not fully understood, it is clear that a variety of chemotactic, cellular, and other environmental signals come into play and within the mesenchymal lineage, for example, mesenchymal stem cells (MSC) cultured in vitro can be induced to differentiate into bone or cartilage in vivo and in vitro, depending upon the tissue environment or the culture medium into which the cells are placed (column 2). Peterson et al also teach that marrow also has the capacity to regenerate bone and other mesenchymal tissue types when implanted in vivo in diffusion chambers (column 1). Peterson et al teach that results of this nature have led to the conclusion that bone marrow contains one or more populations of pluripotent cells, known as stem cells, having the capacity to differentiate into a wide variety of different cell types of the mesenchymal, hematopoietic, and stromal lineages (column 1). Caplan et al, (U.S. Patent No. 5,486,359, published January 23, 1996) teach that isolated human mesenchymal stem cells can differentiate into more than one tissue type (e.g. bone, cartilage, muscle or marrow stroma) (see the Abstract). Wobus (Molecular Aspects of Medicine (2001), 22/3 (149-164) teach that embryonic stem cells are pluripotenet cell line established from undifferentiated embryonic cells characterized

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by nearly unlimited <u>self-renewal and differentiation capacity</u> (see the Abstract). Zhang et al (*The Journal of Clinical Investigation, September 2002, Volume 110, Number 6*) teach that osteogenic fronts of abnormally closing/closed sutures in animals revealed calvarial overgrowth and overlap along with increase osteoblast differentiation and reduced cell proliferation (see the Abstract). Zhang et al teach that anomalies were <u>restricted to calvarial bone</u>, despite generalized, non-tissue-specific over-expression of Nell-1 (see the Abstract). Therefore, the claimed method requires calvarial bone cells.

The cited art has taught that a) osteoblastic cells are believed to be derived from stem cells but it has not been proven, b) mesenchymal stem cells and embryonic stem cells can differentiate into more than one tissue type including one, cartilage, muscle or marrow stroma and c) precise signals that trigger differentiation down a particular path are not fully understood, d) bone marrow contains one or more populations of pluripotent cells, known as stem cells, having the capacity to differentiate into a wide variety of different cell types and e) abnormally closing/closed sutures and overexpression of Nell-1 in animals are restricted to calvarial bone. One skilled in the art would have reason to doubt Applicant's assertion that one could use stem cells and bone marrow cells can used in the claimed method of screening for an agent that modulates bone mineralization comprising contacting the osteogenic cell expressing the NELL-1 gene with a test agent when the cited art has taught that bone marrow and stem cells may not necessarily differentiate into osteogenic cells.

The specification fails to teach or disclose how bone marrow cells or stem cells can be used in the claimed method of screening for agents that modulate bone

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mineralization when these cells may not actually differentiate into osteogenic cells.

There is not teaching or disclosure in the instant specification that shows a method of screening for an agent that modulates bone mineralization using bone marrow or stem cells.

Factors to be considered in determining whether undue experimentation is required are set forth in <u>In re Wands</u> 8 USPQ2d 1400. They include (1) the quantity of experimentation necessary, (2) the amount of direction or guidance presented, (3) the presence or absence of working examples, (4) the nature of the invention, (5) the state of the prior art, (6) the relative skill of those in the art, (7) the predictability or unpredictability of the art and (8) the breadth of the claims.

There is lack of enablement for the use of method for screening an agent that modulates bone mineralization wherein the method comprising contacting an osteogenic cell selected from the group consisting of a stem cell and a bone marrow cell. Therefore, one of skill in the art could not conclude that the Nell-1 gene could be use to screen for agents that modulate bone mineralization using bone marrow cells or stem cells when the instant specification and the cited art teaches that Nell-1 may be associated with intramembraneous bone formation in fetal calvarial osteoblastic cells and bone marrow and stem cell do not necessarily differentiate into these types of cells. It is determined that there are no working examples commensurate with the claims that demonstrate that bone marrow and stem cell can be used in the claimed method. There is limited guidance provided in the specification as to how to use the claimed method since the cited art has taught that the precise signals that trigger differentiation

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down a particular path are not fully understood and there bone marrow and stem cells can differentiate into many other types of cells which are not <u>osteogenic</u> and osteogenic cell are required by the claimed method. The skilled artisan is forced into undue experimentation to practice (make and use) the invention as is broadly claimed.

The following is a quotation of the second paragraph of 35 U.S.C. 112:

The specification shall conclude with one or more claims particularly pointing out and distinctly claiming the subject matter which the applicant regards as his invention.

- 5. Claim 51 is rejected under 35 USC 112 second paragraph for failing to particularly point out and distinctly claim the subject matter which applicant regards as the invention Claim 51 recites "wherein the osteogenic cell is selected from a cell endogenous to a fetal calvarial cell culture. It is unclear as to what Applicant is referring. Clarification is required.
- 6. No claims are allowed.

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Conclusion

7. Any inquiry of the general nature or relating to the status of this general application should be directed to the Group receptionist whose telephone number is (703) 308–0196.

Papers relating to this application may be submitted to Technology Center 1600, Group 1640 by facsimile transmission. The faxing of such papers must conform with the notice published in the Office Gazette, 1096 OG 30 (November 15, 1989). Should applicant wish to FAX a response, the current FAX number for the Group 1600 is (703) 872-9306.

Any inquiry concerning this communication from the examiner should be directed to Vanessa L. Ford, whose telephone number is (571) 272-0857. The examiner can normally be reached on Monday – Friday from 9:00 AM to 6:00 PM. If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Lynette Smith, can be reached at (571) 272-0864.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov./. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).

Vanessa L. Ford Biotechnology Patent Examiner November 9, 2004

PRIMARY EXAMINER

11-12-04